## IN THE CLAIMS

1. (currently amended) A method for forming a high strength hydrogel medical implant comprising:

preparing a polymer solution;

injecting the solution into a mold;

causing said molded solution to gel by physically cross-linking the solution;

adjusting the equilibrium hydrogel crystallinity to insure that the swelling pressure of the hydrogel remains stable after implantation by washing said molded gel in a physiologic solution comprising sodium, carbonate, chloride and potassium ions;

dehydrating the molded gel; and packaging the implant.

- 2. (original) The method as set forth in claim 1 wherein said washing takes place for about one day to twelve weeks.
- 3. (original) The method as set forth in claim 2 wherein the washing takes place from two weeks to twelve weeks.
- 4. (currently amended) The method as set forth in claim 2 wherein the physiologic solution contains is .9% phosphate buffered sodium chloride solution.
- 5. (currently amended) The method as set forth in claim 4 wherein the sodium chloride solution further contains is mixed with a potassium carbonate solution.
- 6. (original) The method as set forth in claim 5 wherein the potassium carbonate solution is between about .025 M and .05 M.
- 7. (original) The method as set forth in claim 6 wherein a .05 M potassium carbonate solution is used for a first portion of the washing and a .025 M potassium carbonate solution is used for a later portion of the washing.

8. (original) The method as set forth in claim 1 wherein the dehydration reduces the water content of the gel to its approximate in vivo equilibrium water content.

- 9. (currently amended) The method as set forth in claim 8 further including irradiating the molded gel after said washing with gamma irradiation.
- 10. (currently amended) The method as set forth in claim 89 wherein said molded gel is hydrated to about 80% water content prior to irradiation.
- 11. (currently amended) The method as set forth in claim 4 wherein said washing in said .9% phosphate buffered sodium chloride saline-solution is for at least-one daytwo weeks.
- 12. (original) The method as set forth in claim 11 wherein said buffered sodium chloride solution includes potassium phosphate.
- 13. (currently amended) A process for treating a hydrogel comprising:

  forming a hydrogel from a polymer solution by physically cross-linking the polymer; and

adjusting the equilibrium hydrogel crystallinity to insure that the swelling pressure of the hydrogel remains stable after implantation by washing the hydrogel in a saline solution including potassium carbonate.

- 14. (original) The process for treating a hydrogel as set forth in claim 13, wherein the saline solution contains between .025 and .05 M potassium carbonate.
- 15. (original) The process for treating a hydrogel as set forth in claim 14, wherein the washing takes place for at least one day.
- 16. (original) The process for treating a hydrogel as set forth in claim 15, wherein the washing takes place for between one day and 12 weeks.

17. (original) The process for treating a hydrogel as set forth in claim 13, wherein the washing solution is heated.

- 18. (original) The process for treating a hydrogel as set forth in claim 17 wherein the solution is heated to 37°C.
- 19. (original) The process for treating a hydrogel as set forth in claim 13, wherein the solution is a .9% phosphate buffered sodium chloride solution with between .025 M and .25 M potassium carbonate added thereto.
- 20. (original) The process as set forth in claim 19 wherein the potassium carbonate added is between .025 M and .05 M.
- 21. (previously presented) The method as set forth in claim 1 wherein the gel formed is semi-crystalline.
- 22. (previously presented) The method as set forth in claim 21 wherein the washing is done for two to twelve weeks in a .9% phosphate buffered sodium chloride solution.
- 23. (previously presented) The method as set forth in claim 22 wherein the solution further contains potassium carbonate.
- 24. (previously presented) The method as set forth in claim 21 wherein the physiologic solution has an ionic charge.
- 25. (previously presented) The method as set forth in claim 1 wherein the polymer is poly (vinyl alcohol).
- 26. (currently amended) The method as set forth in claim 1 wherein the hydrogel is physically cross-linked by a freezing-thawing technique.

27. (currently amended) A method of forming a hydrogel medical implant comprising:

preparing a polymer solution;

physically cross-linking the solution to form a semi-crystalline gel using a freezing-thawing technique;

adjusting the equilibrium hydrogel crystallinity to insure that the swelling pressure of the hydrogel remains stable after implantation by increasing the crystallinity of the gel by washing the gel in a saline solution which further contains potassium carbonate for at least one day.

- 28. (previously presented) The process for treating a hydrogel as set forth in claim 27 wherein the solution is a .9% phosphate buffered sodium chloride solution with between .025 M and .25 M potassium carbonate added thereto.
- 29. (currently amended) The method as set forth in claim 27 further including irradiating the molded gel after said washing with gamma irradiation.
- 30. (previously presented) The method as set forth in claim 27 wherein the polymer is poly (vinyl alcohol).